Reply to Office Action of May 7, 2003

Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

1. (Currently Amended) A method of diagnosing chronic fatigue syndrome in a patient exhibiting symptoms associated with chronic fatigue syndrome, comprising:

evaluating the patient for serologic evidence of EBV and HCMV, further comprising:

obtaining serum from the patient;

measuring the level of EBV IgM antibodies to the VCA in the serum by measuring nonstructural epitopes for incomplete virus multiplication;

measuring the level of EBV antibodies to the total EA in the serum by measuring nonstructural epitopes for incomplete virus multiplication;

measuring the level of HCMV IgM antibodies in the serum by measuring nonstructural epitopes for incomplete virus multiplication;

measuring the level of HCMV IgG antibodies in the serum by measuring nonstructural epitopes for incomplete virus multiplication;

monitoring the patient for T-wave abnormalities;

classifying EBV as the cause of the chronic fatigue syndrome when the measurements show any one of the following: 1) an elevated level of IgM antibodies to the VCA for EBV; and 2) presence of total EA antibodies for EBV, in combination with the absence of IgM antibodies for HCMV and a low level of IgG antibodies for HCMV;

classifying HCMV as the cause of the chronic fatigue syndrome when the measurements show any one of the following: 1) an elevated level of IgM antibodies for HCMV; and 2) an elevated level of IgG antibodies for HCMV, in combination with a low level of IgM antibodies to the VCA for EBV, and the absence of total EA antibodies for EBV; and classifying a combination of EBV and HCMV as the cause of the chronic fatigue syndrome when the measurements show any one of the following: 1) an elevated level of IgM antibodies to the VCA for EBV; and 2) the presence of total EA antibodies for EBV,

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in combination with any of the following: 1) an elevated level of IgM antibodies for HCMV; and 2) an elevated level of IgG antibodies for HCMV.

- 2. (Original) The method of claim 1, wherein the patient's T-waves are monitored through electrocardiographic monitoring.
- 3. (Original) The method of claim 1, wherein the patient's T-waves are monitored through Holter monitoring.
- 4. (Original) The method of claim 1, further comprising the step of conducting a stress multiple gaited acquisition test to check for the presence of an abnormal ventricular dynamics.
- 5. (Original) The method of claim 1, further comprising the step of conducting a myocardial perfusion test to check for coronary artery disease.
- 6. (Original) The method of claim 1, further comprising the step of conducting a cardiac catheterization to determine if a cardiomyopathy exists.
- 7. (Original) The method of claim 1, further comprising the step of conducting an endomyocardial biopsy to check for EBV or HCMV nucleic acids.
- 8. (Original) The method of claim 7, further comprising the step of conducting a polymerase chain reaction study of the biopsy for EBV and HCMV to determine the cause of the chronic fatigue syndrome.
- 9. (Original) The method of claim 7, further comprising the step of conducting in-situ hybridization analysis of the biopsy for EBV and HCMV to determine the cause of the chronic fatigue syndrome.

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10. (Original) A method of diagnosing chronic fatigue syndrome in a patient exhibiting symptoms associated with chronic fatigue syndrome, comprising:

evaluating the patient for serologic evidence of EBV and HCMV, further comprising:

obtaining serum from the patient;

measuring the level of EBV IgM antibodies to the VCA in the

serum by ELISA method;

measuring the level of EBV antibodies to the total EA in the

serum by ELISA method;

measuring the level of HCMV IgM antibodies in the serum by

measuring antigens p52 and CM₂ with the use of a light scattering technique;

measuring the level of HCMV IgG antibodies in the serum by

measuring antigens p52 and CM₂ with the use of a light scattering technique;

monitoring the patient for T-wave abnormalities;

classifying EBV as the cause of the chronic fatigue syndrome when the measurements show any one of the following: 1) an elevated level of IgM antibodies to the VCA for EBV; and 2) presence of total EA antibodies for EBV, in combination with the absence of IgM antibodies for HCMV and a low level of IgG antibodies for HCMV;

classifying HCMV as the cause of the chronic fatigue syndrome when the measurements show any one of the following: 1) an elevated level of IgM antibodies for HCMV; and 2) an elevated level of IgG antibodies for HCMV, in combination with a low level of IgM antibodies to the VCA for EBV, and the absence of total EA antibodies for EBV; and

classifying a combination of EBV and HCMV as the cause of the chronic fatigue syndrome when the measurements show any one of the following: 1) an elevated level of IgM antibodies to the VCA for EBV; and 2) the presence of total EA antibodies for EBV, in combination with any of the following: 1) an elevated level of IgM antibodies for HCMV; and 2) an elevated level of IgG antibodies for HCMV.

11.-23. (Cancelled)